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(54) Titel: HARMFUL FUNGI CONTROL WITH AN ACTIVE SUBSTANCE INHIBITING RESPIRATION BY INHIBITING THE CYTOCHROME COMPLEX III, COMBINED WITH AN AMIDE			
(54) Bezeichnung: SCHADPILZBEKÄMPFUNG MIT DER KOMBINATION EINES DIE ATMUNG AM CYTOCHROM-KOMPLEX III HEMMENDEN WIRKSTOFFES MIT EINEM AMID			
(57) Abstract			
<p>The invention pertains to agents intended for use against harmful fungi and containing as active substances at least one compound which inhibits respiration by inhibiting the cytochrome complex III, and at least one amide compound of formula (II) A-CO-NR¹R² in which A, R¹ and R² have the meanings indicated in the description. The proposed agents are especially suitable for use against <i>Botrytis</i>.</p>			
(57) Zusammenfassung			
<p>Die vorliegende Erfindung betrifft Mittel zur Bekämpfung von Schadpilzen, die als Wirkstoffe wenigstens eine Verbindung, welche die Atmung am Cytochrom-Komplex III hemmt, und wenigstens eine Amidverbindung der Formel (II) A-CO-NR¹R² enthalten, worin A, R¹ und R² die in der Beschreibung angegebenen Bedeutungen besitzen. Die erfindungsgemäßen Mittel sind insbesondere zur Bekämpfung von <i>Botrytis</i> brauchbar.</p>			

O.Z. 0050/46232

Abstract

5 Compositions for controlling harmful fungi which comprise, as active ingredients, at least one compound which inhibits respiration on the cytochrome complex III and at least one amide compound of the formula II

$$10 \quad \text{A} = \text{CO} = \text{NR}^1 \text{R}^2$$

where A, R¹ and R² have the meanings given in the description. The compositions according to the invention can be used, in particular, for controlling botrytis.

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COMPOSITIONS AND METHODS FOR CONTROLLING
HARMFUL FUNGI

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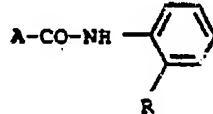
The present invention relates to compositions for controlling harmful fungi and to methods of controlling harmful fungi using such compositions.

10 It is known from the literature that active ingredients which inhibit the cytochrome bc_1 complex (cytochrome complex III) can be employed as fungicides [cf. U. Brandt, U. Haase, H. Schägger, G. von Jagow: "Spezifität und Wirkmechanismus der Strobilurine" (Specificity and mechanism of action of the strobilurins), Dechema monograph Vol. 129, 27-38, VCH Verlagsgesellschaft Weinheim, 1993; J.M. Clough: Natural Product Reports, 1993, 565-574; F. Röhl, H. Sauter: Biochem. Soc. Trans. 22, 635 (1993)].

20 However, when using these active ingredients, it has emerged that their action is only transitory, i.e. new fungal growth was observed after only a short time.

EP-A-545 099 describes anilide compounds of the formula

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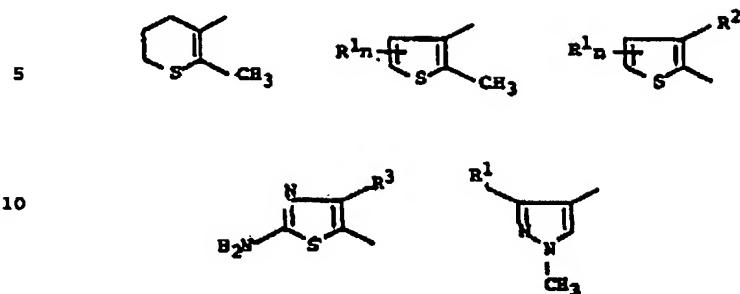
where A is phenyl which is substituted in the 2-position by methyl, trifluoromethyl, chlorine, bromine or iodine, or is one of certain aromatic or non-aromatic heterocyclic radicals which can be unsubstituted or substituted by methyl, chlorine or trifluoromethyl, and R is one of certain aliphatic or cycloaliphatic radicals which can be unsubstituted or substituted by halogen, or is phenyl which is unsubstituted or substituted by C_1-C_4 -alkyl, C_1-C_4 -alkoxy, C_1-C_4 -alkylthio or halogen. These compounds can be used for controlling botrytis.

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EP-A-589 301 describes anilide compounds of the same formula where A is a cyclic radical of the formulae:

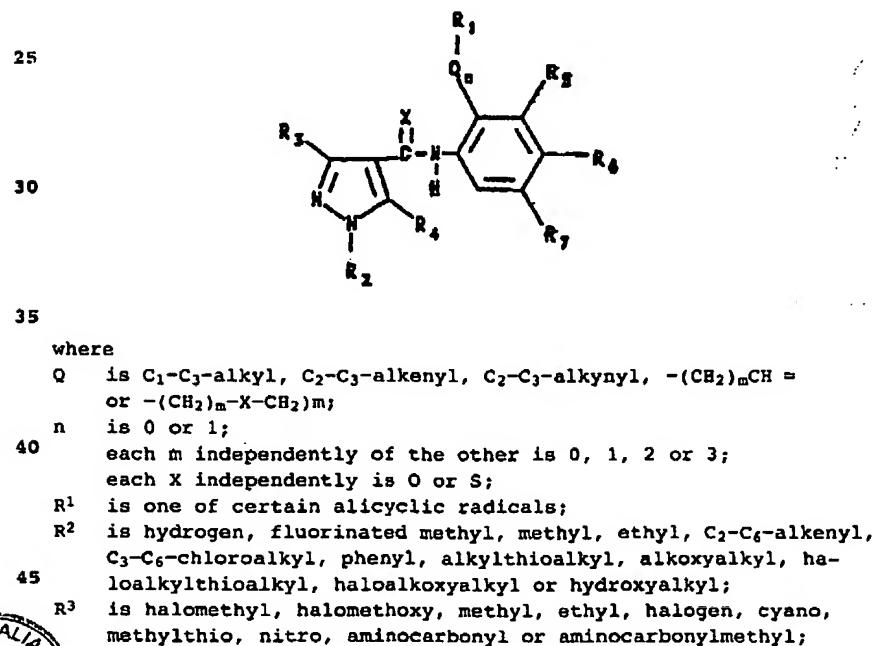
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15 where R^1 is hydrogen or C_1-C_4 -alkyl; R^2 is halogen or C_1-C_4 -alkyl; R^3 is C_1-C_4 -alkyl or C_1-C_4 -haloalkyl; n is 1 or 2; and R has essentially the abovementioned meanings. These compounds can also be used for treating botrytis.

20 WO 93/11117 describes compounds of the formula



R^4 is hydrogen, halogen or methyl;

R^5 , R^6 and R^7 in each case independently of one another are selected from amongst hydrogen, halogen, cyano, C_1 - C_6 -alkyl, C_2 - C_6 -alkenyl, C_2 - C_6 -alkynyl, C_1 - C_4 -alkoxy, C_1 - C_4 -alkylthio, C_3 - C_4 -cycloalkyl and halomethoxy. These compounds are

5 fungicidally active.

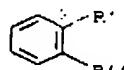
However, it has emerged that the abovementioned anilide compounds do not have a sufficiently broad and satisfactory spectrum of action.

It is an object of the present invention to provide an improved possibility of controlling harmful fungi, in particular botrytis.

10 Surprisingly, we have found that this object is achieved by a composition which comprises an active ingredient which inhibits respiration on cytochrome complex III in combination with an amide compound of the abovementioned type.

The present invention therefore relates to compositions for controlling harmful fungi which comprise, in a solid or liquid carrier,

15 A) at least one active ingredient I, which inhibits respiration on cytochrome complex III, of the formula



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wherein

R' is $-C[CO_2CH_3]=CHOCH_3$, $-C[CO_2CH_3]=NOCH_3$, $-C[CONHCH_3]=NOCH_3$,

$-C[CO_2CH_3]=CHCH_3$, $-C[CO_2CH_3]=CHCH_2CH_3$, $-C[COCH_3]=NOCH_3$,

$-C[COCH_2CH_3]=NOCH_3$, $-N(OCH_3)-CO_2CH_3$, $-N(CH_3)-CO_2CH_3$ or

25 $-N(CH_2CH_3)-CO_2CH_3$,

R'' has one of the following meanings:

a) phenoxyethylene, pyridinyloxyethylene, pyrimidinyloxyethylene or pyrazolylloxyethylene, the aromatic radical being unsubstituted or having 1, 2 or

3 substituents, independently of one another selected from amongst alkyl, halogen,

30 CF_3 , CHF_2 , $-C(CH_3)=NOCH_3$, and phenyl which is unsubstituted or substituted by 1, 2 or 3 halogen atoms and/or alkyl groups;



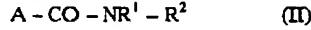
3a

b) phenoxy or pyrimidinyloxy, unsubstituted or substituted by 1, 2 or 3 halogen atoms or a phenoxy radical which is unsubstituted or has a halogen or cyano substituent;

c) $\text{CH}_2\text{ON}=\text{CR}^a\text{R}^b$
5 where R^a is alkyl; and
 R^b is phenyl which is unsubstituted or has 1, 2 or 3 substituents, independently of one another selected from amongst alkyl, halogen, CF_3 and CHF_2 ;

d) $\text{CH}_2\text{ON}=\text{CR}^c\text{R}^d=\text{NOR}^e$
10 where R^c is alkyl;
 R^d is alkyl or phenyl which is unsubstituted or is substituted by 1, 2 or 3 halogen atoms; and R^e is alkyl or alkenyl; and

15 a) at least one active ingredient I, which inhibits respiration on cytochrome complex III, and
b) at least one amide compound of the formula II

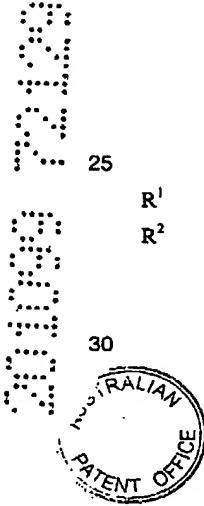


where

20 A is an aryl group or an aromatic or non-aromatic 5- or 6-membered heterocycle which has 1 to 3 hetero atoms selected from amongst O, N and S; it being possible for the aryl group or the heterocycle to be unsubstituted or to have 1, 2 or 3 substituents, independently of one another selected from amongst alkyl, halogen, CHF_2 , CF_3 , alkoxy, haloalkoxy, alkylthio, alkylsulfinyl and alkylsulfonyl;

25 R^f is a hydrogen atom, alkyl or alkoxy;

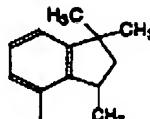
R^g is a phenyl or cycloalkyl group which is unsubstituted or has 1 to 3 substituents, independently of one another selected from amongst phenyl, alkenyl, alkynyl, alkenyloxy, alkynyoxy, cycloalkyl, cycloalkenyl, cycloalkyloxy and cycloalkenyloxy and which can additionally be substituted by 1 or more halogen atoms, it being possible for the



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aliphatic and cycloaliphatic radicals to be partially or fully halogenated and/or for the cycloaliphatic radicals to be substituted by 1 to 3 alkyl groups, and it being possible for the phenyl group, in turn, to have 1 to 5 halogen atoms and/or 1 to 3 substituents, independently of one another selected from amongst alkyl, haloalkyl, alkoxy, haloalkoxy, alkylthio and haloalkylthio, and it being possible for the amidic phenyl group to be fused to a saturated 5-membered ring which is unsubstituted or substituted by one or more alkyl groups and/or can have one hetero atom selected from amongst O and S, with the exception of the compound of the formula II where

15 A is 2-chloropyridin-3-yl, R¹ is H and R² is



25 The compositions according to the invention have a synergistic action and are therefore especially suitable for controlling harmful fungi, in particular *botrytis*. It is assumed that this is based on the fact that, when respiration on the cytochrome complex III is inhibited, the fungus utilizes a secondary route of 30 alternative respiration, so that the fungi are not destroyed completely. This would mean that the amide compounds of the formula II inhibit the alternative respiration. It must therefore be assumed that the combination of the two active ingredients inhibits respiration via the cytochrome complex III and also the alternative respiration, so that the fungi are destroyed completely. 35

Halogen within the scope of the present invention is fluorine, chlorine, bromine and iodine, in particular fluorine, chlorine and bromine.

40 The term "alkyl" encompasses straight-chain or branched alkyl groups. They are preferably straight-chain or branched C₁-C₁₂-alkyl and, in particular, C₁-C₆-alkyl groups. Examples of alkyl groups are alkyl such as, in particular, methyl, ethyl, propyl, 1-methylethyl, butyl, 1-methylpropyl, 2-methylpropyl 45 1,1-dimethylethyl, n-pentyl, 1-methylbutyl, 2-methylbutyl, 3-methylbutyl, 1,2-dimethylpropyl, 1,1-dimethylpropyl, 2,2-dimethylpropyl, 1-ethylpropyl, n-hexyl, 1-methylpentyl, 2-methylpentyl.



3-methylpentyl, 4-methylpentyl, 1,2-dimethylbutyl, 1,3-dimethylbutyl, 2,3-dimethylbutyl, 1,1-dimethylbutyl, 2,2-dimethylbutyl, 3,3-dimethylbutyl, 1,1,2-trimethylpropyl, 1,2,2-trimethylpropyl, 1-ethylbutyl, 2-ethylbutyl, 1-ethyl-2-methylpropyl,
 5 n-heptyl, 1-methylhexyl, 1-ethylpentyl, 2-ethylpentyl, 1-propylbutyl, octyl, decyl, dodecyl.

Haloalkyl is an alkyl group as defined above which is partially or fully halogenated with one or more halogen atoms, in particular fluorine and chlorine. There are preferably 1 to 3 halogen atoms present, the difluoromethane [sic] or the trifluoromethyl group being especially preferred.

15 What has been said above about the alkyl group and haloalkyl group applies analogously to the alkyl and haloalkyl group in alkoxy, haloalkoxy, alkylthio, haloalkylthio, alkylsulfinyl and alkylsulfonyl.

20 The alkenyl group encompasses straight-chain and branched alkenyl groups. Preferably, they are straight-chain or branched C₃-C₁₂-alkenyl groups, in particular C₃-C₆-alkenyl groups. Examples of alkenyl groups are 2-propenyl, 2-butenyl, 3-butenyl, 1-methyl-2-propenyl, 2-methyl-2-propenyl, 2-pentenyl, 3-pentenyl, 4-pentenyl, 1-methyl-2-butenyl, 2-methyl-2-butenyl,
 25 3-methyl-2-butenyl, 1-methyl-3-butenyl, 2-methyl-3-butenyl, 3-methyl-3-butenyl, 1,1-dimethyl-2-propenyl, 1,2-dimethyl-2-propenyl, 1-ethyl-2-propenyl, 2-hexenyl, 3-hexenyl, 4-hexenyl, 5-hexenyl, 1-methyl-2-pentenyl, 2-methyl-2-pentenyl, 3-methyl-2-pentenyl, 4-methyl-2-pentenyl, 1-methyl-3-pentenyl,
 30 2-methyl-3-pentenyl, 3-methyl-3-pentenyl, 4-methyl-3-pentenyl, 1-methyl-4-pentenyl, 2-methyl-4-pentenyl, 3-methyl-4-pentenyl, 4-methyl-4-pentenyl, 1,1-dimethyl-2-butenyl, 1,1-dimethyl-3-butenyl, 1,1-dimethyl-3-butyl, 1,2-dimethyl-3-butenyl, 1,2-dimethyl-2-butenyl, 1,2-dimethyl-3-butenyl, 1,3-dimethyl-2-butenyl, 1,3-dimethyl-3-butenyl,
 35 2,2-dimethyl-3-butenyl, 2,3-dimethyl-2-butenyl, 2,3-dimethyl-3-butenyl, 1-ethyl-2-butenyl, 1-ethyl-3-butenyl, 2-ethyl-2-butenyl, 2-ethyl-3-butenyl, 1,1,2-trimethyl-2-propenyl, 1-ethyl-1-methyl-2-propenyl and 1-ethyl-2-methyl-2-propenyl, in particular 2-propenyl, 2-butenyl, 3-methyl-2-butenyl and
 40 3-methyl-2-pentenyl.

The alkenyl group can be partially or fully halogenated with one or more halogen atoms, in particular fluorine and chlorine. Preferably, it has 1 to 3 halogen atoms.



The alkynyl group encompasses straight-chain and branched alkynyl groups. They are preferably straight-chain and branched C₃-C₁₂-alkynyl groups, in particular C₃-C₆-alkynyl groups.

Examples of alkynyl groups are 2-propynyl, 2-butynyl, 3-butynyl,

5 1-methyl-2-propynyl, 2-pentynyl, 3-pentynyl, 4-pentynyl, 1-methyl-3-butynyl, 2-methyl-3-butynyl, 1-methyl-2-butynyl, 1,1-dimethyl-2-propynyl, 1-ethyl-2-propynyl, 2-hexynyl, 3-hexynyl, 4-alkynyl, 5-hexynyl, 1-methyl-2-pentynyl, 1-methyl-3-pentynyl, 1-methyl-4-pentynyl, 2-methyl-3-pentynyl, 2-methyl-4-penty-
10 nyl, 3-methyl-4-pentynyl, 4-methyl-2-pentynyl, 1,2-dimethyl-2-butynyl, 1,1-dimethyl-3-butynyl, 1,2-dimethyl-3-butynyl, 2,2-dimethyl-3-butynyl, 1-ethyl-2-butynyl, 1-ethyl-3-butynyl, 2-ethyl-3-butynyl and 1-ethyl-1-methyl-2-propynyl.

15 What has been said above about the alkenyl group and its halogen substituents, and about the alkynyl group, applies analogously to alkenyloxy and alkynyloxy.

The cycloalkyl group is preferably a C₃-C₆-cycloalkyl group, such as cyclopropyl, cyclobutyl, cyclopentyl or cyclohexyl. If the cycloalkyl group is substituted, it preferably has 1 to 3 C₁-C₄-alkyl radicals as substituents.

20 Cycloalkenyl is preferably a C₄-C₆-cycloalkenyl group, such as cyclobutenyl, cyclopentenyl or cyclohexenyl. If the cycloalkenyl group is substituted, it preferably has 1 to 3 C₁-C₄-alkyl radicals as substituents.

25 A cycloalkoxy group is preferably a C₅-C₆-cycloalkoxy group, such as cyclopentyloxy or cyclohexyloxy. If the cycloalkoxy group is substituted, it preferably has 1 to 3 C₁-C₄-alkyl radicals as substituents.

30 The cycloalkenyloxy group is preferably a C₅-C₆-cycloalkenyloxy group, such as cyclopentyloxy or cyclohexyloxy. If the cycloalkenyloxy group is substituted, it preferably has 1 to 3 C₁-C₄-alkyl radicals as substituents.

Aryl is preferably phenyl.

35 Hetaryl is preferably a 5- or 6-membered aromatic heterocycle which has 1, 2 or 3 hetero atoms, independently of one another selected from amongst N, O and S. It is, in particular, pyridinyl, pyrimidinyl, thiazolyl, pyrazolyl, oxazolyl, isoxazolyl, iso-thiazolyl, imidazolyl, pyrrolyl, furanyl, thieryl or triazolyl.

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Heterocyclyl is preferably a 5- or 6-membered saturated or unsaturated heterocycle which has 1, 2 or 3 hetero atoms, independently of one another selected from amongst N, O and S. It is, in particular, a dihydro, tetrahydro or hexahydro derivative of the radicals mentioned under "hetaryl". Pyrrolidinyl, tetrahydrofuryl, imidazolidinyl, pyrazolidinyl, oxazolidinyl, isoxazolidinyl, thiazolidinyl, isothiazolidinyl, piperidinyl or morpholinyl are preferred.

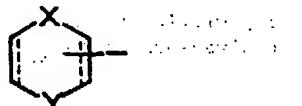
10 If A in formula II is a phenyl group, it can have one, two or three of the abovementioned substituents in any position. Preferably, these substituents, independently of one another, are selected from amongst alkyl,

15 difluoromethyl, trifluoromethyl and halogen, in particular chlorine, bromine and iodine. Especially preferably, the phenyl group has a substituent in the 2-position.

20 If A is a 5-membered heterocycle, this is, in particular, a furyl, thiazolyl, pyrazolyl, imidazolyl, oxazolyl, thienyl, triazolyl or thiadiazolyl radical or a corresponding dihydro or tetrahydro derivative thereof. A thiazolyl or pyrazolyl radical is preferred.

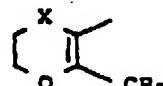
25 If A is a 6-membered heterocycle, this is, in particular, a pyridyl radical or a radical of the formula:

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35 The 6-membered aromatic heterocycle is especially preferably a pyridyl radical, in particular a 3-pyridyl radical, or a radical of the formula

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(A3)

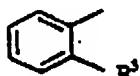
45 where X is CH_2 , S, SO or SO_2 .



The abovementioned heterocyclic radicals may be unsubstituted or have 1, 2 or 3 of the abovementioned substituents, these substituents, preferably, independently of one another being selected from amongst alkyl, halogen, difluoromethyl or trifluoromethyl.

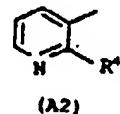
A especially preferably is a radical of the formulae:

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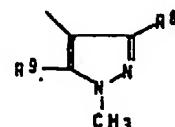
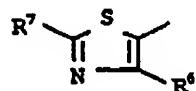
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(A1)



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(A5)

(A7)

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where R³, R⁴, R⁶, R⁷, R⁸ and R⁹ independently of one another are hydrogen, alkyl, in particular methyl, halogen, in particular chlorine, CHF_2 or CF_3 .

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The radical R¹ in formula II is preferably a hydrogen atom.

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The radical R² in formula II is preferably a phenyl radical. R² preferably has at least one substituent, particularly preferably in the 2-position. The substituent (or substituents) is/are preferably selected from amongst alkyl, cycloalkyl, cycloalkenyl, halogen or phenyl.

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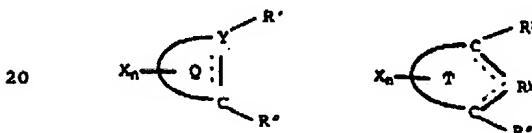
The substituents of the radical R² can, in turn, also be substituted. The aliphatic or cycloaliphatic substituents can be partially or fully halogenated, in particular fluorinated or chlorinated. They preferably have 1, 2 or 3 fluorine or chlorine atoms. If the substituent of the radical R² is a phenyl group, it



can preferably be substituted by 1 to 3 halogen atoms, in particular chlorine atoms and/or by a radical which is preferably selected from amongst alkyl and alkoxy. Especially preferably, the phenyl group is substituted in the *p*-position with a halogen atom, i.e. the especially preferred substituent of the radical R² is a *p*-halogen-substituted phenyl radical. The radical R² can also be fused to a saturated 5-membered ring, it being possible for this ring, in turn, to have 1 to 3 alkyl substituents. In this case, R² is, for example, indanyl, thiaindanyl and oxaindanyl. Preferred are indanyl and 2-oxaindanyl, which are bonded to the nitrogen atom in particular via the 4-position.

The active ingredient I is preferably a compound of the formula IA or IB:

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where is a double or single bond;

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R' is -C[CO₂CH₃]=CHOCH₃, -C[CO₂CH₃]=NOCH₃, -C[CONHCH₃]=NOCH₃, -C[CO₂CH₃]=CHCH₃, -C[CO₂CH₃]=CHCH₂CH₃, -C[COCH₃]=NOCH₃, -C[COCH₂CH₃]=NOCH₃, -N(OCH₃)-CO₂CH₃, -N(CH₃)-CO₂CH₃ or -N(CH₂CH₃)-CO₂CH₃

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R'' is an organic radical which is bonded directly or via an oxy, mercapto, amino, or alkylamino group, or

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together with a group X and the ring Q or T to which they are bonded an unsubstituted or substituted bicyclic, partially or fully unsaturated system which, besides carbon ring members, may contain 1, 2 or 3 hetero atoms, independently selected from amongst oxygen, sulfur and nitrogen;

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R^X is -OC[CO₂CH₃]=CHOCH₃, -OC[CO₂CH₃]=CHCH₃, -OC[CO₂CH₃]=CHCH₂CH₃, -SC[CO₂CH₃]=CHOCH₃, -SC[CO₂CH₃]=CHCH₃, -SC[CO₂CH₃]=CHCH₂CH₃, -N(CH₃)C[CO₂CH₃]=CHOCH₃, -N(CH₃)C[CO₂CH₃]=NOCH₃, -CH₂C[CO₂CH₃]=CHOCH₃, -CH₂C[CO₂CH₃]=NOCH₃ or -CH₂C[CONHCH₃]=NOCH₃;



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10
R' is oxygen, sulfur, $=\text{CH}-$ or $=\text{N}-$;

n is 0, 1, 2 or 3, it being possible for the radicals X to be identical or different if $n > 1$;

5
X is cyano, nitro, halogen, alkyl, haloalkyl, alkoxy, haloalkoxy, alkylthio, or,

10 if $n > 1$, a $\text{C}_3\text{-C}_5$ -alkylene, $\text{C}_3\text{-C}_5$ -alkenylene, oxy- $\text{C}_2\text{-C}_4$ -alkylene, oxy- $\text{C}_1\text{-C}_3$ -alkyleneoxy, oxy- $\text{C}_2\text{-C}_4$ -alkenylene, oxy- $\text{C}_2\text{-C}_4$ -alkenyleneoxy or butadienediyl group bonded to two adjacent C atoms of the phenyl ring, it being possible for these chains, in turn, to have attached to them one to three radicals, independently of one another selected from amongst halogen, alkyl, haloalkyl, alkoxy, haloalkoxy and alkylthio;

15
Y is $=\text{C}-$ or $=\text{N}-$;

20 Q is phenyl, pyrrolyl, thiienyl, furyl, pyrazolyl, imidazolyl, oxazolyl, isoxazolyl, thiazolyl, thiadiazolyl, triazolyl, pyridinyl, 2-pyridonyl, pyrimidinyl or triazinyl; and

25 T is phenyl, oxazolyl, thiazolyl, thiadiazolyl, oxadiazolyl, pyridinyl, pyrimidinyl or triazinyl.

25 The substituent R" is, in particular, an alkyl, alkenyl, alkynyl, aryl, hetaryl, arylalkyl, hetarylalkyl, arylalkenyl, hetarylalkenyl, arylalkynyl or hetarylalkynyl radical which is uninterrupted or interrupted by one or more groups selected from amongst 30 O, S, SO , SO_2 , NR (R = H or alkyl), CO, COO , OCO , CONH , NHCO and NHCNH , or a radical of the formula $\text{CH}_2\text{ON} = \text{CR}^\alpha\text{CR}^\beta$ or $\text{CH}_2\text{ON} = \text{CR}^\gamma\text{CR}^\delta = \text{NOR}'$ defined below. These radicals are unsubstituted or have one or more (preferably 1, 2 or 3) substituents, independently of one another selected from amongst alkyl, alkoxy, halogen, cyano, haloalkyl (in particular CF_3 and CHF_2), hetaryl and aryl. Hetaryl and aryl can, in turn, have 1, 2 or 3 substituents which, independently of one another, are selected from amongst 35 halogen, haloalkyl (in particular CF_3 and CHF_2), phenyl, CN, phenoxy, alkyl, alkoxy and haloalkoxy.

40 Such compounds and their preparation are described in the literature given in Tables I.1 to I.8 below. Compounds not described therein can be prepared by similar methods.

45 In a preferred embodiment, the compositions according to the invention comprise a compound of the formula IA or IB where R" is aryloxy, hetaryloxy, aryloxymethylene, hetaryloxymethylene, arylethenylene or hetarethylene, these radicals being unsub-



stituted or having 1, 2 or 3 substituents, independently of one another selected from amongst alkyl, halogen, CF_3 , CHF_2 , CN , alkoxy and phenyl which, in turn, can have 1, 2 or 3 substituents, independently of one another selected from amongst halogen, CF_3 , CHF_2 , phenyl, CN , phenoxy, alkyl, alkoxy and haloalkoxy;

or R'' is



where R^a is alkyl;

15 R^b is phenyl, pyridyl or pyrimidyl, unsubstituted or having 1, 2 or 3 substituents independently of one another selected from amongst alkyl, alkoxy, halogen, haloalkoxy, CF_3 and CHF_2 ;

R^c is alkyl, alkoxy, halogen, haloalkyl or hydrogen;

20 R^d [sic] is hydrogen, cyano, halogen, alkyl, alkoxy, alkylthio, alkylamino, dialkylamino, alkenyl, alkenyloxy, alkennythio, alkenylamino, N -alkenyl- N -alkylamino, alkynyl, alkynyloxy, alkynylthio, alkynylamino, N -alkynyl- N -alkylamino, it being possible for the hydrocarbon radicals of these groups to be 25 partially or fully halogenated and/or to have attached to them 1, 2 or 3 radicals, independently selected from amongst cyano, nitro, hydroxyl, alkoxy, haloalkoxy, alkoxy carbonyl, alkylthio, alkylamino, dialkylamino, alkenyloxy, cycloalkyl, cycloalkyloxy, heterocyclyl, heterocyclxyloxy, aryl, aryloxy, arylalkoxy, hetaryl, hetaryloxy and hetarylalkoxy, it being 30 possible for the cyclic radicals, in turn, to be partially or fully halogenated and/or to have attached to them 1, 2 or 3 groups, independently selected from amongst cyano, nitro, hydroxyl, alkyl, haloalkyl, cycloalkyl, alkoxy, haloalkoxy, alkoxy carbonyl, alkylthio, alkylamino, dialkylamino, alkenyl 35 and alkenyloxy;

or

40 is cycloalkyl, cycloalkyloxy, cycloalkylthio, cycloalkylamino, N -cycloalkyl- N -alkylamino, heterocyclyl, heterocyclxyloxy, heterocyclylthio, heterocyclylamino, N -heterocyclyl- N -alkylamino, aryl, aryloxy, arylthio, arylamino, N -aryl- N -alkylamino, hetaryl, hetaryloxy, hetarylthio, hetarylamino or 45 N -hetaryl- N -alkylamino, it being possible for the cyclic radicals to be partially or fully halogenated and/or to have attached to them 1, 2 or 3 groups, independently selected from



12

amongst cyano, nitro, hydroxyl, alkyl, haloalkyl, cycloalkyl, alkoxy, haloalkoxy, alkoxy carbonyl, alkylthio, alkylamino, dialkylamino, alkenyl, alkenyloxy, benzyl, benzyloxy, aryl, aryloxy, hetaryl and hetaryloxy, it being possible for the aromatic radicals, in turn, to be partially or fully halogenated and/or to have attached to them 1, 2 or 3 of the following groups: cyano, alkyl, haloalkyl, alkoxy, nitro;

5 R^c is alkyl, alkenyl or alkynyl, it being possible for these
10 groups to be partially or fully halogenated and/or to have attached to them 1, 2 or 3 of the following radicals: cyano, alkoxy, cycloalkyl.

Especially preferred are compounds of the formula IA or IB where
15 R^c has one of the following meanings:

a) phenoxy methylene, pyridinyloxy methylene, pyrimidinyloxy methylene or pyrazolyl oxy methylene, the aromatic radical being unsubstituted or having 1, 2 or 3 substituents, independently
20 of one another selected from amongst alkyl, halogen, CF₃, CHF₂, -C(CH₃)=NOCH₃, and phenyl which is unsubstituted or substituted by 1, 2 or 3 halogen atoms and/or alkyl groups;

b) phenoxy or pyrimidinyloxy, unsubstituted or substituted by 1, 2 or 3 halogen atoms or a phenoxy radical which is unsubstituted or has a halogen or cyano substituent;

c) phenylethylene or pyrazolylethylene, it being possible for the phenyl or pyrazolyl radical to be unsubstituted or to have 1, 2 or 3 substituents, independently of one another selected from amongst halogen, CF₃, CHF₂ and phenyl.

d) CH₂ON=CR^aR^b

35 where R^a is alkyl; and

R^b is phenyl which is unsubstituted or has 1, 2 or 3 substituents, independently of one another selected from amongst alkyl, halogen, CF₃ and CHF₂, or is pyrimidinyl which is unsubstituted or is substituted by 1 or 2 alkoxy radicals;

e) CH₂ON=CR^cCR^d=NOR^e

where R^c is alkyl, alkoxy or halogen;



13

R^{δ} is alkyl, cyano, halogen, alkoxy, alkenyl or phenyl which is unsubstituted or is substituted by 1, 2 or 3 halogen atoms; and

5 R^{δ} is alkyl.

Particularly preferred are compounds of the formula IA where Q is phenyl and n is 0.

10 Especially suitable active ingredients I are compiled in the Tables which follow.

Table I.1A

15 Compounds of the formula IA where Q is phenyl, R' is $-\text{C}(\text{CO}_2\text{CH}_3)=\text{CHOCH}_3$, n has a value of 0, R^{δ} is unsubstituted or substituted (het)aryloxymethylene, the unsubstituted or substituted (het)aryl group having the following meaning

No.	Unsubst. or subst. (het)aryl	Literature
I.1A-1	2-CH ₃ -C ₆ H ₄	EP-A 226 917
I.1A-2	2,5-(CH ₃) ₂ -C ₆ H ₃	EP-A 226 917
I.1A-3	2-CH ₃ , 4-C(CH ₃)=NOCH ₃ -C ₆ H ₃	EP-A 386 561
I.1A-4	2-CH ₂ CH ₂ CH ₃ , 6-CF ₃ -pyrimidin-4-yl	EP-A 407 873
I.1A-5	2,4-(CH ₃) ₂ -C ₆ H ₃	EP-A 226 917

Table I.1B:

30 Compounds of the formula IA where R' is $-\text{C}(\text{CO}_2\text{CH}_3)=\text{CHOCH}_3$, Q is phenyl, n has a value of 0, R^{δ} is unsubstituted or substituted (het)aryloxy, the unsubstituted or substituted (het)aryl group having the following meaning

35

No.	Unsubst. or subst. (het)aryl	Literature
I.1B-1	C ₆ H ₅	EP-A 178 826
I.1B-2	6-[2-CN-C ₆ H ₄ -O]-pyrimidin-4-yl	EP-A 382 375

40

Table I.1C:

Compounds of the formula IA where R' is $-\text{C}(\text{CO}_2\text{CH}_3)=\text{CHOCH}_3$, Q is phenyl, n has a value of 0, R^{δ} is unsubstituted or substituted (het)arylethenylene, the unsubstituted or substituted (het)aryl group having the following meaning



No.	Unsubst. or subst. (het)aryl	Literature
I.1C-1	1-(2,4-Cl ₂ -C ₆ H ₃), 5-CF ₃ -pyrazol-4-yl	EP-A 528 245
I.1C-2	1-(4-Cl-C ₆ H ₄)-pyrazol-4-yl	EP-A 378 755
I.1C-3	3-CF ₃ -C ₆ H ₄	EP-A 203 606
I.1C-4	3-Cl-C ₆ H ₄	EP-A 203 606
I.1C-5	4-C ₆ H ₅ -C ₆ H ₄	EP-A 203 606

10 Table I.1D:

15 Compounds of the formula IA where Q is phenyl, R' is
-C(CO₂CH₃)=CHOCH₃, n has a value of 0, R" is CH₂ON=CR^aR^b, R^a and R^b
having the following meanings

No.	R ^a	R ^b	Literature
I.1D-1	CH ₃	4-Cl-C ₆ H ₄	EP-A 370 629
I.1D-2	CH ₃	3-CF ₃ -C ₆ H ₄	EP-A 370 629
I.1D-3	CH ₃	4-OCH ₂ CH ₃ -pyrimidin-2-yl	WO-A 92/18,487

Table I.1E:

20 Compounds of the formula IA where Q is phenyl, R' is
-C(CO₂CH₃)=CHOCH₃, n has a value of 0, R" is CH₂ON=CR^yCR^b=NOR^e, R^y,
R^b and R^e having the following meanings

No.	R ^y	R ^b	R ^e	Literature
I.1E-1	CH ₃	CH ₃	CH ₃	WO-A 95/21153
I.1E-2	CH ₃	CH ₃	CH ₂ CH ₃	WO-A 95/21153
I.1E-3	CH ₃	C ₆ H ₅	CH ₃	WO-A 95/21153
I.1E-4	CH ₃	C ₆ H ₅	CH ₂ CH ₃	WO-A 95/21153
I.1E-5	CH ₃	4-Cl-C ₆ H ₄	CH ₃	WO-A 95/21153
I.1E-6	CH ₃	4-Cl-C ₆ H ₄	CH ₂ CH ₃	WO-A 95/21153

Table I.2A:

25 Compounds of the formula IA where Q is phenyl, R' is
-C(CO₂CH₃)=NOCH₃, n has a value of 0, R" is unsubstituted or sub-
stituted (het)aryloxymethylene, the unsubstituted or substituted
(het)aryl group having the following meaning



No.	Unsubst. or subst. (het)aryl	Literature
5	I.2A-1 2-CH ₃ -C ₆ H ₄	EP-A 253 213
	I.2A-2 2,5-(CH ₃) ₂ -C ₆ H ₃	EP-A 400 417
	I.2A-3 2,4-(CH ₃) ₂ -C ₆ H ₃	EP-A 400 417
	I.2A-4 2,3,5-(CH ₃) ₃ -C ₆ H ₂	EP-A 400 417
	I.2A-5 2-Cl, 5-CH ₃ -C ₆ H ₃	EP-A 400 417
	I.2A-6 2-CH ₃ , 4-C(CH ₃)=NOCH ₃ -C ₆ H ₃	EP-A 386 561

10 Table I.2B:

Compounds of the formula IA where Q is phenyl, R' is -C(CO₂CH₃)=NOCH₃, n has a value of 0, R" is unsubstituted or substituted (het)aryloxy, the unsubstituted or substituted (het)aryl group having the following meaning

No.	Unsubst. or subst. (het)aryl	Literature
20	I.2B-1 C ₆ H ₅	EP-A 253 213
	I.2B-2 6-[2-CN-C ₆ H ₄ -O]-pyrimidin-4-yl	EP-A 468 684

Table I.2C:

Compounds of the formula IA where Q is phenyl, R' is -C(CO₂CH₃)=NOCH₃, n has a value of 0, R" is CH₂ON=CR^aR^b, R^a and R^b having the following meanings

No.	R ^a	R ^b	Literature
30	CH ₃	4-Cl-C ₆ H ₄	EP-A 463 488
	CH ₃	3-Cl-C ₆ H ₄	EP-A 463 488
	CH ₃	4-CF ₃ -C ₆ H ₄	EP-A 463 488
	CH ₃	3-CF ₃ -C ₆ H ₄	EP-A 463 488
35	CH ₃	4-CH ₃ -C ₆ H ₄	EP-A 463 488
	CH ₃	4-OCH ₂ CH ₃ -pyrimidin-2-yl	EP-A 472 300
	CH ₃	3,5-Cl ₂ -C ₆ H ₃	EP-A 463 488

40 Table I.2D:

Compounds of the formula IA where Q is phenyl, R' is -C(CO₂CH₃)=NOCH₃, n has a value of 0, R" is CH₂ON=CR^aR^b=NOR^c, R^a, R^b and R^c having the following meanings



No.	R'	R ^b	R"	Literature
5	I.2D-1	CH ₃	CH ₃	WO-A 95/21153
	I.2D-2	CH ₃	CH ₂ CH ₃	WO-A 95/21153
	I.2D-3	CH ₃	C ₆ H ₅	WO-A 95/21153
	I.2D-4	CH ₃	C ₆ H ₅	WO-A 95/21153
	I.2D-5	CH ₃	4-Cl-C ₆ H ₄	WO-A 95/21153
	I.2D-6	CH ₃	4-Cl-C ₆ H ₄	WO-A 95/21153

10 Table I.3A:

Compounds of the formula IA where Q is phenyl, R' is -C(CONHCH₃)=NOCH₃, n has a value of 0, R" is unsubstituted or substituted (het)aryloxymethylene, the unsubstituted or substituted (het)aryl group having the following meaning

No.	Unsubst. or subst. (het)aryl	Literature
I.3A-1	2-CH ₃ -C ₆ H ₄	EP-A 477 631
I.3A-2	2,5-(CH ₃) ₂ -C ₆ H ₃	EP-A 477 631
I.3A-3	2,4-(CH ₃) ₂ -C ₆ H ₃	EP-A 477 631
I.3A-4	2,3,5-(CH ₃) ₃ -C ₆ H ₂	EP-A 477 631
I.3A-5	2-CH ₃ , 4-C(CH ₃)=NOCH ₃ -C ₆ H ₃	EP-A 579 124
I.3A-6	1-[4-Cl-C ₆ H ₄]-pyrazol-3-yl	WO-A 94/19331
I.3A-7	1-[2,4-Cl ₂ -C ₆ H ₃]-pyrazol-3-yl	WO-A 94/19331

20 Table I.3B:

30 Compounds of the formula IA where Q is phenyl, R' is -C(CONHCH₃)=NOCH₃, n has a value of 0, R" is unsubstituted or substituted (het)aryloxy, the unsubstituted or substituted (het)aryl group having the following meaning

No.	Unsubst. or subst. (het)aryl	Literature
I.3B-1	C ₆ H ₅	EP-A 398 692
I.3B-2	6-[2-CN-C ₆ H ₄ -O]-pyrimidin-4-yl	GB-A 2 253 624

40 Table I.3C:

Compounds of the formula IA where Q is phenyl, R' is -C(CONHCH₃)=NOCH₃, n has a value of 0, R" is unsubstituted or substituted (het)arylethylene, the unsubstituted or substituted (het)aryl group having the following meaning



No.	Unsubst. or subst. (het)aryl	Literature
I.3C-1	1-[2,4-Cl ₂ -C ₆ H ₃], 5-CF ₃ -pyrazol-4-yl	DE-A 44 23 615.8

5 Table I.3D:

Compounds of the formula IA where Q is phenyl, R' is -C(CONHCH₃)=NOCH₃, n has a value of 0, R" is CH₂ON=CR^aR^b, R^a and 10 R^b having the following meanings

No.	R ^a	R ^b	Literature
I.3D-1	CH ₃	4-Cl-C ₆ H ₄	EP-A 463 488
I.3D-2	CH ₃	3-Cl ₃ -C ₆ H ₄	EP-A 463 488
I.3D-3	CH ₃	4-CF ₃ -C ₆ H ₄	EP-A 585 751
I.3D-4	CH ₃	3-CF ₃ -C ₆ H ₄	EP-A 585 751
I.3D-5	CH ₃	4-CH ₃ -C ₆ H ₄	EP-A 463 488
I.3D-6	CH ₃	3,5-Cl ₂ -C ₆ H ₃	EP-A 463 488
I.3D-7	CH ₃	2-OCH ₂ CH ₃ -pyrimidin-2-yl	WO-A 92/13,830

15 Table I.3E:

20 Compounds of the formula IA where Q is phenyl, R' is -C(CONHCH₃)=NOCH₃, n has a value of 0, R" is CH₂ON=CR^aR^b, R^a, R^b and R^c having the following meanings

No.	R ^a	R ^b	R ^c	Literature
I.3E-1	CH ₃	CH ₃	CH ₃	WO-A 95/21154
I.3E-2	CH ₃	CH ₃	CH ₂ CH ₃	WO-A 95/21154
I.3E-3	CH ₃	C ₆ H ₅	CH ₃	WO-A 95/21154
I.3E-4	CH ₃	C ₆ H ₅	CH ₂ CH ₃	WO-A 95/21154
I.3E-5	CH ₃	4-Cl-C ₆ H ₄	CH ₃	WO-A 95/21154
I.3E-6	CH ₃	4-Cl-C ₆ H ₄	CH ₂ CH ₃	WO-A 95/21154
I.3E-7	CH ₃	4-F-C ₆ H ₄	CH ₃	WO-A 95/21154

25 Table I.4A:

Compounds of the formula IA where Q is phenyl, R' is -C(CO₂CH₃)=CHCH₃, n has a value of 0, R" is unsubstituted or substituted (het)aryloxymethylene, the unsubstituted or substituted (het)aryl group having the following meaning



NO.	Unsubst. or subst. (het)aryl	Literature
I.4A-1	2-CH ₃ -C ₆ H ₄	EP-A 280 185
I.4A-2	2,5-(CH ₃) ₂ -C ₆ H ₃	EP-A 513 580
5 I.4A-3	2,4-(CH ₃) ₂ -C ₆ H ₃	EP-A 513 580
I.4A-4	2,3,5-(CH ₃) ₃ -C ₆ H ₂	EP-A 513 580
I.4A-5	2-Cl, 5-CH ₃ -C ₆ H ₃	EP-A 513 580
I.4A-6	2-CH ₃ , 4-C(CH ₃)=NOCH ₃ -C ₆ H ₃	EP-A 513 580
10 I.4A-7	1-[4-Cl-C ₆ H ₄]-pyrazol-3-yl	DE-A 44 15 483.6

Table I.4B:

15 Compounds of the formula IA where Q is phenyl, R' is -C(CO₂CH₃)=CHCH₃, n has the value of 0, R" is unsubstituted or substituted (het)aryloxy, the unsubstituted or substituted (het)aryl group having the following meaning

NO.	Unsubst. or subst. (het)aryl	Literature
I.4B-1	C ₆ H ₅	EP-A 513 580

Table I.4C:

20 25 Compounds of the formula IA where Q is phenyl, R' is -C(CO₂CH₃)=CHCH₃, n has a value of 0, R" is CH₂ON=CR^YCR^δ=NOR^ε, R^Y, R^δ and R^ε having the following meanings

NO.	R ^Y	R ^δ	R ^ε	Literature
I.4C-1	CH ₃	CH ₃	CH ₃	WO-A 95/21153
I.4C-2	CH ₃	CH ₃	CH ₂ CH ₃	WO-A 95/21153
I.4C-3	CH ₃	C ₆ H ₅	CH ₃	WO-A 95/21153
30 I.4C-4	CH ₃	C ₆ H ₅	CH ₂ CH ₃	WO-A 95/21153
I.4C-5	CH ₃	4-Cl-C ₆ H ₄	CH ₃	WO-A 95/21153
35 I.4C-6	CH ₃	4-Cl-C ₆ H ₄	CH ₂ CH ₃	WO-A 95/21153

Table I.5A:

40 Compounds of the formula IA where Q is phenyl, R' is -C(CO₂CH₃)=CHCH₂CH₃, n has a value of 0, R" is unsubstituted or substituted (het)aryloxymethylene, the unsubstituted or substituted (het)aryl group having the following meaning



M/36140

No.	Unsubst. or subst. (het)aryl	Literature
I.5A-1	2-CH ₃ -C ₆ H ₄	EP-A 513 580
I.5A-2	2,5-(CH ₃) ₂ -C ₆ H ₃	EP-A 513 580
I.5A-3	2,4-(CH ₃) ₂ -C ₆ H ₃	EP-A 513 580
I.5A-4	2,3,5-(CH ₃) ₃ -C ₆ H ₂	EP-A 513 580
I.5A-5	2-Cl, 5-CH ₃ -C ₆ H ₃	EP-A 513 580
I.5A-6	2-CH ₃ , 4-C(CH ₃)=NOCH ₃ -C ₆ H ₃	EP-A 513 580

10 Table I.5B:

Compounds of the formula IA where Q is phenyl, R' is -C(CO₂CH₃)=CHCH₂CH₃, n has a value of 0, R'' is unsubstituted or substituted (het)aryloxy, the unsubstituted or substituted (het)aryl group having the following meaning

No.	Unsubst. or subst. (het)aryl	Literature
I.5B-1	C ₆ H ₅	EP-A 513 580

20 Table I.5C:

Compounds of the formula IA where Q is phenyl, R' is -C(CO₂CH₃)=CHCH₂CH₃, n has a value of 0, R'' is CH₂ON=CRYCR^δ=NOR^ε, R^γ, R^δ and R^ε having the following meanings

No.	R ^γ	R ^δ	R ^ε	Literature
I.5C-1	CH ₃	CH ₃	CH ₃	WO-A 957 21153
I.5C-2	CH ₃	CH ₃	CH ₂ CH ₃	WO-A 95/21153
I.5C-3	CH ₃	C ₆ H ₅	CH ₃	WO-A 95/21153
I.5C-4	CH ₃	C ₆ H ₅	CH ₂ CH ₃	WO-A 95/21153
I.5C-5	CH ₃	4-Cl-C ₆ H ₄	CH ₃	WO-A 95/21153
I.5C-6	CH ₃	4-Cl-C ₆ H ₄	CH ₂ CH ₃	WO-A 95/21153

30 Table I.6A

Compounds of the formula IA where Q is phenyl, R' is -C(COCH₃)=NOCH₃, n has a value of 0, R'' is unsubstituted or substituted (het)aryloxymethylene, the unsubstituted or substituted (het)aryl group having the following meaning

No.	Unsubst. or subst. (het)aryl	Literature
I.6A-1	2-CH ₃ -C ₆ H ₄	EP-A 498 188
I.6A-2	2,5-(CH ₃) ₂ -C ₆ H ₃	EP-A 498 188



I.6A-3	2,4-(CH ₃) ₂ -C ₆ H ₃	EP-A 498 188
I.6A-4	2,3,5-(CH ₃) ₃ -C ₆ H ₂	EP-A 498 188
I.6A-5	2-CH ₃ , 4-C(CH ₃)=NOCH ₃ -C ₆ H ₃	EP-A 498 188

5

Table I.6B:

Compounds of the formula IA where Q is phenyl, R' is -C(COCH₃)=NOCH₃, n has a value of 0, R" is unsubstituted or substituted (het)aryloxy, the unsubstituted or substituted (het)aryl group having the following meaning

No.	Unsubst. or subst. (het)aryl	Literature
I.6B-1	C ₆ H ₅	EP-A 498 188
I.6B-2	6-[2-CN-C ₆ H ₄ -O]-pyrimidin-4-yl	EP-A 498 188

Table I.7A:

Compounds of the formula IA where Q is phenyl, R' is -C(COCH₂CH₃)=NOCH₃, n has a value of 0, R" is unsubstituted or substituted (het)aryloxymethylene, the unsubstituted or substituted (het)aryl group having the following meaning

No.	Unsubst. or subst. (het)aryl	Literature
I.7A-1	2-CH ₃ -C ₆ H ₄	EP-A 498 188
I.7A-2	2,5-(CH ₃) ₂ -C ₆ H ₃	EP-A 498 188
I.7A-3	2,4-(CH ₃) ₂ -C ₆ H ₃	EP-A 498 188
I.7A-4	2,3,5-(CH ₃) ₃ -C ₆ H ₂	EP-A 498 188
I.7A-5	2-CH ₃ , 4-C(CH ₃)=NOCH ₃ -C ₆ H ₃	EP-A 498 188

Table I.7B:

Compounds of the formula IA where Q is phenyl, R' is -C(COCH₂CH₃)=NOCH₃, n has a value of 0, R" is unsubstituted or substituted (het)aryloxy, the unsubstituted or substituted (het)aryl group having the following meaning

No.	Unsubst. or subst. (het)aryl	Literature
I.7B-1	C ₆ H ₅	EP-A 498 188
I.7B-2	6-[2-CN-C ₆ H ₄ -O]-pyrimidin-4-yl	EP-A 498 188



Table I.8A:

Compounds of the formula IA where Q is phenyl, R' is
 5 $-N(OCH_3)-CO_2CH_3$, n has a value of 0, R'' is unsubstituted or sub-
 substituted (het)aryloxymethylene, the unsubstituted or substituted
 (het)aryl group having the following meaning

NO.	Unsubst. or subst. (het)aryl	Literature
10	I.8A-1 2-CH ₃ -C ₆ H ₄	WO-A 93/15,046
	I.8A-2 2,5-(CH ₃) ₂ -C ₆ H ₃	WO-A 93/15,046
	I.8A-3 2,4-(CH ₃) ₂ -C ₆ H ₃	WO-A 93/15,046
15	I.8A-4 2,3,5-(CH ₃) ₃ -C ₆ H ₂	WO-A 93/15,046
	I.8A-5 2-Cl, 5-CH ₃ -C ₆ H ₃	WO-A 93/15,046
	I.8A-6 2-CH ₃ , 4-C[CH ₃]=NOCH ₃ -C ₆ H ₃	WO-A 93/15,046
	I.8A-7 2-CH ₃ , 4-C[CH ₃]=NOCH ₂ CH ₃ -C ₆ H ₃	WO-A 93/15,046
20	I.8A-8 2-CH ₃ , 4-C[CH ₂ CH ₃]=NOCH ₃ -C ₆ H ₃	WO-A 93/15,046
	I.8A-9 2-CH ₃ , 4-C[CH ₂ CH ₃]=NOCH ₂ CH ₃ -C ₆ H ₃	WO-A 93/15,046
	I.8A-10 1-[4-Cl-C ₆ H ₄]-pyrazol-3-yl	DE-A 44 23 612.3

Table I.8B:

25 Compounds of the formula IA where Q is phenyl, R' is
 $-N(OCH_3)-CO_2CH_3$, n has a value of 0, R'' is $CH_2ON=CR^aR^b$, R^a and R^b
 having the following meanings

NO.	R ^a	R ^b	Literature
30	I.8B-1 CH ₃	3,5-Cl ₂ -C ₆ H ₃	WO-A 93/15,046

In a further preferred embodiment, the composition according to
 the invention comprises, as the amide compound, a compound of the
 35 formula II where A has the following meanings:
 phenyl, pyridyl, dihydropyranyl, dihydrooxathiinyl, dihydrooxa-
 thiinyl oxide, dihydrooxathiinyl dioxide, furyl, thiazolyl, pyra-
 zolyl or oxazolyl, it being possible for these groups to have 1,
 2 or 3 substituents, independently of one another selected from
 40 amongst alkyl, halogen, difluoromethyl and trifluoromethyl.

In a further preferred embodiment, A represents:
 pyridin-3-yl which is unsubstituted or substituted in the 2-posi-
 tion by halogen, methyl, difluoromethyl, trifluoromethyl, me-
 thoxy, methylthio, methylsulfinyl or methylsulfonyl;



22

phenyl which is unsubstituted or substituted in the 2-position by methyl, trifluoromethyl, chlorine, bromine or iodine;

5 2-methyl-5,6-dihydropyran-3-yl;

2-methyl-5,6-dihydro-1,4-oxathiin-3-yl or the 4-oxide or 4,4-dioxide thereof;

10 2-methyl-furan-3-yl which is unsubstituted or substituted in the 4- and/or 5-position by methyl;

thiazol-5-yl which is unsubstituted or substituted in the 2- and/or 4-position by methyl, chlorine, difluoromethyl or trifluoromethyl;

15 2-methylpyrazol-4-yl which is unsubstituted or substituted in the 2- and/or 5-position by methyl, chlorine, difluoromethyl or trifluoromethyl;

20 1-methylpyrazol-4-yl which is unsubstituted or substituted in the 3- and/or 5-position by methyl, chlorine, difluoromethyl or trifluoromethyl; or

25 oxazol-5-yl which is unsubstituted or substituted in the 2- and/or 4-position by methyl or chlorine.

30 In a further preferred embodiment, the compositions according to the invention comprise, as the amide compound, a compound of the formula II where R² is a phenyl group which is unsubstituted or substituted by 1, 2 or 3 of the abovementioned substituents.

35 In a further preferred embodiment, the compositions according to the invention comprise, as the amide compound, a compound of the formula II where R² is a phenyl group which has one of the following substituents in the 2-position:

C₃-C₆-alkyl, C₃-C₆-cycloalkenyl, C₃-C₆-cycloalkyloxy, C₃-C₆-cycloalkenylloxy, it being possible for these groups to be substituted by 1, 2 or 3 C₁-C₄-alkyl groups,

40 phenyl which is substituted by 1 to 5 halogen atoms and/or 1 to 3 groups independently of one another selected from amongst C₁-C₄-alkyl, C₁-C₄-haloalkyl, C₁-C₄-alkoxy, C₁-C₄-haloalkoxy, C₁-C₄-alkylthio and C₁-C₄-haloalkylthio,

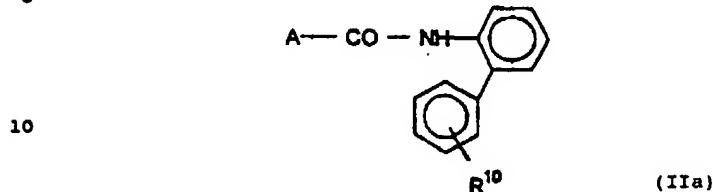
45 indanyl or oxaindanyl, unsubstituted or substituted by 1, 2 or 3 C₁-C₄-alkyl groups.



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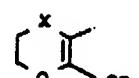
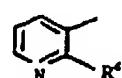
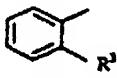
In a further preferred embodiment, the compositions according to the invention comprise, as the amide compound, a compound of the formula IIIa,

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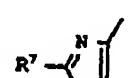
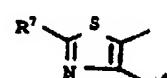


where

15 A is



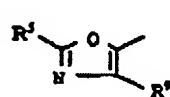
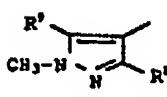
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X is methylene, sulfur, sulfinyl or sulfonyl (SO_2),

R³ is methyl, difluoromethyl, trifluoromethyl, chlorine, bromine or iodine,

R⁴ is trifluoromethyl or chlorine,

R⁵ is hydrogen or methyl,

R⁶ is methyl, difluoromethyl, trifluoromethyl or chlorine,



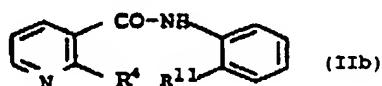
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R⁷ is hydrogen, methyl or chlorine,
R⁸ is methyl, difluoromethyl or trifluoromethyl,
R⁹ is hydrogen, methyl, difluoromethyl, trifluoromethyl or
chlorine, and
5 R¹⁰ is C₁-C₄-alkyl, C₁-C₄-alkoxy, C₁-C₄-alkylthio or halogen.

In an especially preferred embodiment, the compositions comprise,
as the amide compound, a compound of the formula IIb

10

15



(IIb)

20 where

R⁴ is halogen and
R¹¹ is phenyl which is substituted by halogen.

25 Suitable amide compounds are mentioned in EP-A-545 099 and
589 301, which are herewith referred to in their entirety.

The preparation of the amide compounds of the formula I is dis-
closed, for example, in EP-A-545 099 or 589 301 or can be carried
30 out by similar processes.

To obtain the synergistic effect, the active ingredients are
employed in a weight ratio in a range of from 20:1 to 1:20, in
particular 10:1 to 1:10.

35 The invention also relates to a method of controlling harmful
fungi, which comprises treating the fungi, their environment, or
the materials, plants, seeds, soils, areas or spaces to be pro-
tected against fungal infection, with a composition as defined
40 above, it being possible for the active ingredients to be applied
simultaneously, i.e. jointly or separately, or in succession.

The compositions according to the invention can be applied, for
example, in the form of directly sprayable solutions, powders,
45 suspensions, also highly-concentrated aqueous, oily or other sus-
pensions or dispersions, emulsions, oil dispersions, pastes,
dusts, materials for spreading or granules by means of spraying,
atomizing, dusting, spreading or pouring. The use forms depend on



the intended purposes; in any case, they should guarantee the finest possible distribution of the active ingredients according to the invention.

5 Normally, the plants are sprayed or dusted with the active ingredients, or the seeds of the plants are treated with the active ingredients.

The formulations are prepared in a known manner, eg. by extending 10 the active ingredient with solvents and/or carriers, if desired using emulsifiers and dispersants, it also being possible to use other organic solvents as auxiliary solvents if water is used as the diluent. Auxiliaries which are suitable are essentially: solvents such as aromatics (eg. xylene), chlorinated aromatics (eg. 15 chlorobenzenes), paraffins (eg. mineral oil fractions), alcohols (eg. methanol, butanol), ketones (eg. cyclohexanone), amines (eg. ethanolamine, dimethylformamide) and water; carriers such as ground natural minerals (eg. kaolins, clays, talc, chalk) and ground synthetic minerals (eg. highly-dispersed silica, silicates); emulsifiers, such as non-ionic and anionic emulsifiers 20 (eg. polyoxyethylene fatty alcohol ethers, alkylsulfonates and arylsulfonates) and dispersants such as lignin-sulfite waste liquors and methylcellulose.

25 Suitable surfactants are the alkali metal, alkaline earth metal and ammonium salts of aromatic sulfonic acids, eg. ligno-, phenol-, naphthalene- and dibutylnaphthalenesulfonic acid, and of fatty acids, of alkyl- and alkylarylsulfonates, of alkyl sulfates, lauryl ether sulfates and fatty alcohol sulfates; and salts of sulfated hexa-, hepta- and octadecanols, or else fatty 30 alcohol glycol ethers, condensates of sulfonated naphthalene and its derivatives with formaldehyde, condensates of naphthalene or of the naphthalene sulfonic acids with phenol and formaldehyde, polyoxyethylene octylphenyl ether, ethoxylated isoctyl-, octyl- or nonylphenol, alkylphenol [sic] polyglycol ethers, tributylphenyl polyglycol ether, alkylaryl polyether alcohols, isotridecyl alcohol, fatty alcohol/ethylene oxide condensates, ethoxylated castor oil, polyoxyethylene alkyl ethers or polyoxypropylene 35 [sic], lauryl alcohol polyglycol ether acetate, sorbitol esters, lignin-sulfite waste liquors or methylcellulose.

40 Powders, materials for spreading and dusts can be prepared by mixing or concomitantly grinding the active substances with a solid carrier.

45 Granules, eg. coated granules, impregnated granules and homogeneous granules, can be prepared by binding the active ingredients to solid carriers. Solid carriers are mineral earths, such as



silica gel, silicas, silica gels [sic], silicates, talc, kaolin, limestone, lime, chalk, bole, loess, clay, dolomite, diatomaceous earth, calcium sulfate, magnesium sulfate, magnesium oxide, ground synthetic materials, fertilizers, such as ammonium sulfate, ammonium phosphate, ammonium nitrate, ureas, and products of vegetable origin, such as cereal meal, tree bark meal, wood meal and nutshell meal, cellulose powders or other solid carriers.

10 Examples of such preparations which comprise the active ingredients in a weight ratio of 1:1 are:

I. a solution of 90 parts by weight of the active ingredients and 10 parts by weight of N-methylpyrrolidone which is 15 suitable for use in the form of microdrops;

II. a mixture of 20 parts by weight of the active ingredients, 80 parts by weight of xylene, 10 parts by weight of the adduct of 8 to 10 mol of ethylene oxide to 1 mol of oleic acid 20 N-monoethanolamide, 5 parts by weight of calcium dodecylbenzenesulfonate, 3 parts by weight of the adduct of 40 mol of ethylene oxide to 1 mol of castor oil; a dispersion is obtained by finely distributing the solution in water;

III. an aqueous dispersion of 20 parts by weight of the active ingredients, 40 parts by weight of cyclohexanone, 30 parts 25 by weight of isobutanol, 20 parts by weight of the adduct of 40 mol of ethylene oxide to 1 mol of castor oil;

IV. an aqueous dispersion of 20 parts by weight of the active ingredients, 25 parts by weight of cyclohexanol, 65 parts by 30 weight of a mineral oil fraction of boiling point 210 to 280°C and 10 parts by weight of the adduct of 40 mol of ethylene oxide to 1 mol of castor oil;

V. a mixture, ground in a hammer mill, of 80 parts by 35 weight of the active ingredients, 3 parts by weight of sodium diisobutylnaphthalene-1-sulfonate, 10 parts by weight of the sodium salt of a lignosulfonic acid from a sulfite waste liquor and 7 parts by weight of pulverulent silica gel; a spray mixture is obtained by finely distributing the mixture in water;

VI. an intimate mixture of 3 parts by weight of the active ingredients and 97 parts by weight of finely divided kaolin; this dust comprises 3% by weight of active ingredient;

VII. an intimate mixture of 30 parts by weight of the active ingredients, 92 parts by weight of pulverulent silica gel 45 and 8 parts by weight of paraffin oil which had been sprayed onto



the surface of this silica gel; this preparation imparts good adhesive properties to the active ingredient;

5 VIII. a stable aqueous dispersion of 40 parts by weight of the active ingredients, 10 parts by weight of the sodium salt of a phenolsulfonic acid/urea/formaldehyde condensate, 2 parts by weight of silica gel and 48 parts by weight of water; this dispersion can be diluted further;

10 IX. a stable oily dispersion of 20 parts by weight of the active ingredients, 2 parts by weight of calcium dodecylbenzenesulfonate, 8 parts by weight of fatty alcohol polyglycol ether, 20 parts by weight of the sodium salt of a phenolsulfonic acid/urea/formaldehyde condensate and 88 parts by weight of a paraffin mineral oil.

15 The compositions according to the invention are distinguished by an outstanding activity against a broad spectrum of phytopathogenic fungi, in particular against botrytis. Some of them act 20 systemically (ie. they can be taken up by the treated plants without losing their activity and, if appropriate, translocated within the plant) and can be employed as foliar- and soil-acting fungicides.

25 They are particularly important for controlling a large number of fungi in a variety of crop plants such as wheat, rye, barley, oats, rice, maize, grass, cotton, soybeans, coffee, sugar cane, grapevine, fruit species, ornamentals and vegetable species such as cucumbers, beans and cucurbits, and on the seeds of these plants.

30 The compositions are applied by treating the fungi, or the seeds, plants, materials or the soil to be protected against fungal infection, with a fungicidally active amount of the active ingredients.

35 Application is effected before or after infection of the materials, plants or seeds with the fungi.

40 Specifically, the compositions are suitable for controlling the following plant diseases:

Erysiphe graminis (powdery mildew) in cereals,
Erysiphe cichoracearum and Sphaerotheca fuliginea in cucurbits,
Podosphaera leucotricha in apples,
45 Uncinula necator in grapevines,
Venturia inaequalis (scab) in apples,
Helminthosporium species in cereals,



Septoria nodorum in wheat,
Botrytis cinerea (gray mold) in strawberries and grapevines,
Cercospora arachidicola in groundnuts,
Pseudocercospora herpotrichoides in wheat and barley,
5 Pyricularia oryzae in rice,
Fusarium and Verticillium species in a variety of plants, Alternaria species in vegetables and fruit,
Monilinia species in fruit,
Sclerotinia species in oilseed rape and vegetables.

10 The use against botrytis is preferred.

The compositions can also be employed in the protection of material (protection of wood), for example against Paecilomyces variotii.

In general, the fungicidal compositions comprise from 0.1 to 95, preferably 0.5 to 90, % by weight of active ingredient.

20 Depending on the nature of the desired effect, the rates of application are from 0.02 to 3 kg of active ingredient per ha.
In the treatment of seeds, amounts of from 0.001 to 50 g, preferably 0.01 to 10 g, of active ingredient are generally required per kilogram of seed.

25 In the use form as fungicides, the compositions according to the invention can also comprise other active ingredients, eg. herbicides, insecticides, growth regulators, fungicides or else fertilizers.

30 In many cases, a mixture with fungicides results in a widened fungicidal spectrum of action.

The following list of fungicides together with which the compounds according to the invention can be used is intended to 35 illustrate the possible combinations, but not to impose any limitations:

sulfur,
dithiocarbonates and their derivatives, such as
40 iron(III) dimethyldithiocarbamate,
zinc dimethyldithiocarbamate,
zinc ethylenebisdithiocarbamate,
manganese ethylenebisdithiocarbamate,
manganese zinc ethylenediaminebisdithiocarbamate,
45 tetrathiomolybdate [sic],
ammonia complex of zinc (N,N-ethylenebisdithiocarbamate),
ammonia complex of zinc (N,N'-propylenebisdithiocarbamate),



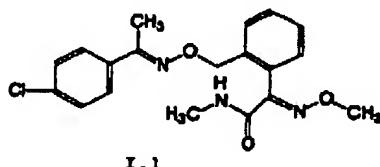
zinc (N,N'-propylenebisdithiocarbamate),
 N,N'-polypropylenebis(thiocarbamoyl)disulfide,
 nitro derivatives, such as
 dinitro(1-methylheptyl)phenyl crotonate,
 5 2-sec-butyl-4,6-dinitrophenyl 3,3-dimethylacrylate,
 2-sec-butyl-4,6-dinitrophenylisopropyl carbonate,
 diisopropyl 5-nitroisophthalate:
 heterocyclic substances, such as
 10 2-heptadecyl-2-imidazoline acetate,
 2,4-dichloro-6-(o-chloroanilino)-s-triazine,
 O,O-diethyl phthalimidophosphonothioate,
 5-amino-1-βbis(dimethylamino)phosphinyl)0-3-phenyl-1,2,4-triazole
 [sic],
 2,3-dicyano-1,4-dithianthraquinone,
 15 2-thio-1,3-dithiolo β 4,5-βquinoxaline [sic],
 1-(butylcarbamoyl)-2-benzimidazolecarbamate,
 2-methoxycarbonylaminobenzimidazole,
 2-(furyl-(2))benzimidazole,
 2-(thiazolyl-(4))benzimidazole,
 20 N-(1,1,2,2-tetrachloroethylthio)tetrahydropthalimide,
 N-trichloromethylthiotetrahydropthalimide,
 N-trichloromethylthiophthalimide,
 N-dichlorofluoromethylthio-N',N'-dimethyl-N-phenylsulfuramide,
 5-ethoxy-3-trichloromethyl-1,2,3-thiadiazole,
 25 2-thiocyanatomethylthiobenzothiazole,
 1,4-dichloro-2,5-dimethoxybenzene,
 4-(2-chlorophenylhydrazono)-3-methyl-5-isoxazolone,
 pyridine-2-thiol 1-oxide,
 8-hydroxyquinoline and its copper salt,
 2,3-dihydro-5-carboxanilido-6-methyl-1,4-oxathine,
 30 2,3-dihydro-5-carboxanilido-6-methyl-1,4-oxathine 4,4-dioxide,
 2-methyl-5,6-dihydro-4H-pyran-3-carboxanilide,
 2-methylfuran-3-carboxanilide,
 2,5-dimethylfuran-3-carboxanilide,
 2,4,5-trimethylfuran-3-carboxanilide,
 35 N-cyclohexyl-2,5-dimethylfuran-3-carboxamide,
 N-cyclohexyl-N-methoxy-2,5-dimethylfuran-3-carboxamide,
 2-methylbenzanilide,
 2-iodobenzanilide,
 N-formyl-N-morpholine-2,2,2-trichloroethyl acetate,
 40 piperazine-1,4-diylbis(1-(2,2,2-trichloroethyl)formamide [sic],
 1-(3,4-dichloroanilino)-1-formylamino-2,2,2-trichloroethane,
 2,6-dimethyl-N-tridecylmorpholine and its salts,
 2,6-dimethyl-N-cyclododecylmorpholine and its salts,
 N-[3-(p-tert-butylphenyl)-2-methylpropyl]-cis-2,6-dimethylmorpho-
 45 line,
 3-(p-tert-butylphenyl)-2-methylpropyl]piperidine,



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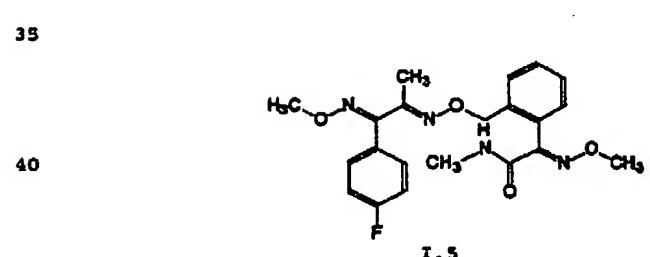
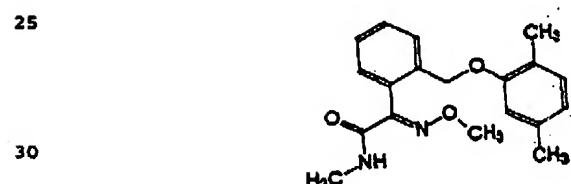
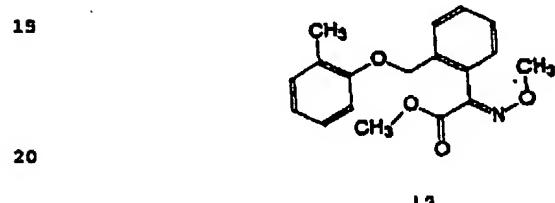
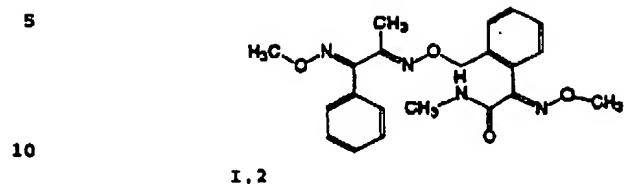
1-[2-(2,4-dichlorophenyl)-4-ethyl-1,3-dioxolan-2-yl-ethyl]-1H-1,2,4-triazole
 1-[2-(2,4-dichlorophenyl)-4-n-propyl]-1,3-dioxolan-2-yl-ethyl]-1H-1,2,4-triazole
 5 N-(n-propyl)-N-(2,4,6-trichlorophenoxyethyl)-N'-imidazolyurea,
 1-(4-chlorophenoxy)-3,3-dimethyl-1-(1H-1,2,4-triazol-1-yl)-2-butane,
 1-(4-chlorophenyl)-3,3-dimethyl-1-(1H-1,2,4-triazol-1-yl)-2-butanol,
 10 α-(2-chlorophenyl)-α-(4-chlorophenyl)-5-pyrimidinemethanol,
 5-butyl-2-dimethylamino-4-hydroxy-6-methylpyrimidine,
 bis(p-chlorophenyl)-3-pyridinemethanol,
 1,2-bis(3-ethoxycarbonyl-2-thioureido)benzene,
 1,2-bis(3-methoxycarbonyl-2-thioureido)benzene,
 15 and a variety of fungicides, such as
 dodecylguanidine acetate,
 3-(3-(3,5-dimethyl)-2-oxycyclohexyl)-2-hydroxyethyl]-glutarimide,
 hexachlorobenzene,
 20 DL-methyl-N-(2,6-dimethylphenyl)-N-(2-furoyl)alaninate,
 DL-N-(2,6-dimethylphenyl)-N-(2'-methoxyacetyl)alanine methyl ester,
 N-(2,6-dimethylphenyl)-N-chloroacetyl-D,L-2-aminobutyrolactone,
 DL-N-(2,6-dimethylphenyl)-N-(phenylacetyl)alanine methyl ester,
 25 5-methyl-5-vinyl-3-(3,5-dichlorophenyl)-2,4-dioxo-1,3-oxazolidine,
 3-(3,5-dichlorophenyl)-(5-methyl-5-methoxymethyl)-1,3-oxazolidine-2,4-dione (sic),
 3-(3,5-dichlorophenyl)-1-isopropylcarbamoylhydantoin,
 N-(3,5-dichlorophenyl)-1,2-dimethylcyclopropane-1,2-dicarboximide,
 30 2-cyano-[N-(ethylaminocarbonyl)-2-methoximino]acetamide,
 1-[2-(2,4-dichlorophenyl)pentyl]-1H-1,2,4-triazole,
 2,4-difluoro-α-(1H-1,2,4-triazolyl-1-methyl)benzhydryl alcohol,
 N-(3-chloro-2,6-dinitro-4-trifluoromethylphenyl)-5-trifluoromethyl-35
 1-(bis(4-fluorophenyl)methylsilyl)methyl-1H-1,2,4-triazole.

The synergistic action of the compositions according to the invention is illustrated with the aid of the Use Examples which 40 follow, the active ingredients I which are employed being the compounds of the formulae I.1 to I.5



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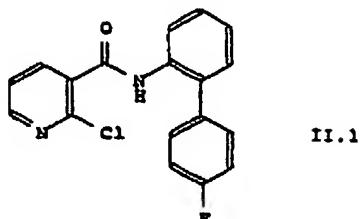


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and the amide compounds employed being the compounds of the formulae II.1 and II.2

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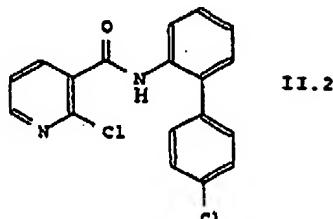
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II.1

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II.2

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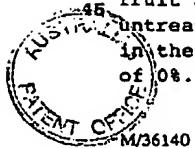
Use Example 1

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Activity against Botrytis cinerea on bell pepper fruits

Disks of green bell pepper fruits were sprayed to drip point with an aqueous preparation of the active ingredient comprising 80% of 35 active ingredient and 20% of emulsifier in the dry matter. 2 hours after the spray coating had dried on, the fruit disks were inoculated with a spore suspension of *Botrytis cinerea* comprising 1.7×10^6 spores per ml of a 2% strength Biomaltz solution. The inoculated fruit disks were subsequently inoculated [sic] for 4 40 days in humid chambers at 18°C. The botrytis development on the infected fruit disks was then evaluated visually (disease level 100%).

The visually determined values for the percentage of infected fruit area were converted into efficacies expressed in % of the 45 untreated control. An efficacy of 0 is the same disease level as in the untreated control, an efficacy of 100 is a disease level of 0%. The expected efficacies for active ingredient combinations



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were determined using Colby's formula (Colby, S. R. (Calculating synergistic and antagonistic responses of herbicide Combinations", Weeds, 15, pp. 20 - 22, 1967) and compared with the observed efficacies. The results are shown in Table 1 below.

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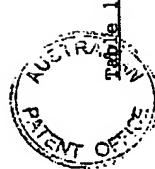
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Active ingredient	Active ingredient concentration in ppm		Efficacy in % of the control	
	I.1 - I.3	II.1	Observed	Calculated*
Control (untreated)	—	—	0	—
II.1	15	50	50	50
	7.5	20		
I.1	5	25		
	2.5	15		
I.2	5	20		
	2.5	10		
I.3	5	5		
	2.5	0		
I.1 + II.1	5	15	99	62
	2.5	7.5	97	32
I.2 + II.1	5	15	96	55
	2.5	15	86	52
I.3 + II.1	5	7.5	60	20
	2.5			

* calculated using Colby's formula

Table 1



The test results show that the efficacy observed in all mixing ratios exceeds the additive efficacy calculated beforehand using Colby's formula, ie. a synergistic effect is present.

5

Use Example 2

Activity against Botrytis cinerea

10 After 4 - 5 leaves had developed properly, bell pepper seedlings cv. "Neusiedler Ideal Elite" were sprayed to drip point with aqueous suspensions comprising 80% of active ingredient and 20% of emulsifier in the dry matter. After the spray coating had dried on, the plants were sprayed with a conidia suspension of 15 the fungus Botrytis cinerea and placed into a chamber at 22 - 24°C and high atmospheric humidity. After 5 days, the untreated control plants had developed such an extent of the disease that the leaf necroses formed covered most of the leaves (disease level 96%).

20 The visually determined values for the percentage of infected leaf area were converted into efficacies expressed in % of the untreated control. An efficacy of 0 is the same disease level as in the untreated control, an efficacy of 100 is a disease level of 0%. The expected efficacies for active ingredient combinations 25 were determined using Colby's formula (Colby, S. R. (Calculating synergistic and antagonistic responses of herbicide Combinations", Weeds, 15, pp. 20 - 22, 1967) and compared with the observed efficacies. The results are shown in Table 2 below.

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35

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Table 2

5	Active ingredient	Active ingredient concentration in ppm		Efficacy in % of the control	
		I.1 - I.3	II.1	Observed	Calculated*
10	Control (untreated)			0	
	II.2	--	24 6 1.5	87 79 28	--
15	I.1	2 0.5	--	0 0	--
	I.2	2 0.5	--	0 0	--
	I.3	2 0.5	--	0 0	--
20	I.4	2 0.5	--	0 0	--
	I.5	2 0.5	--	0 0	--
25	I.1 + II.2	2 0.5	6 1.5	100 100	87 79
	I.2 + II.2	2 0.5	6 1.5	100 99	87 79
	I.3 + II.2	2 0.5	6 1.5	100 99	87 79
30	I.4 + II.2	2 0.5	6 1.5	100 100	87 79
	I.5 + II.2	2 0.5	6 1.5	99 90	87 79

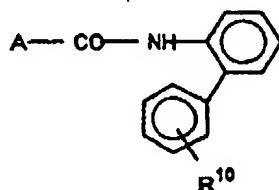
35 * calculated using Colby's formula



The test results show that the efficacy observed in all mixing 5 ratios exceeds the additive efficacy calculated beforehand using Colby's formula, ie. a synergistic effect is present.

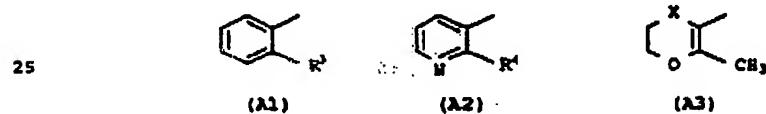
Similar results are obtained when the amide compound employed is one of the compounds of the formula Ia

10



where A is

20



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which are mentioned in Table 3 below, or another of the individual compounds mentioned in EP-A-545 099 and 589 301 and/or one of the individual compounds mentioned in the above Tables I.1 to



No.	A	R ³	R ⁴	R ⁵	R ⁶	R ⁷	R ⁸	R ⁹	R ¹⁰	X	Physical data (°C)
3.1	A ₁	CH ₃	-	-	-	-	-	-	2-F	-	-
3.2	A ₁	CH ₃	-	-	-	-	-	-	4-F	-	-
3.3	A ₁	CF ₃	-	-	-	-	-	-	2-F	-	-
3.4	A ₁	CF ₃	-	-	-	-	-	-	4-F	-	-
3.5	A ₂	-	C ₁	-	-	-	-	-	2-F	-	-
3.6	A ₂	-	C ₁	-	-	-	-	-	2-CH ₃	-	71 - 73
3.7	A ₂	-	C ₁	-	-	-	-	-	2-C ₁	-	-
3.8	A ₂	-	C ₁	-	-	-	-	-	2-0CH ₃	-	-
3.9	A ₂	-	C ₁	-	-	-	-	-	3-F	-	-
3.10	A ₂	-	C ₁	-	-	-	-	-	3-C ₁	-	95 - 98
3.11	A ₂	-	C ₁	-	-	-	-	-	3-CH ₃	-	-
3.12	A ₂	-	C ₁	-	-	-	-	-	3-0CH ₃	-	-
3.13	A ₂	-	C ₁	-	-	-	-	-	3-O(C ₃ H ₇)	-	-
3.14	A ₂	-	C ₁	-	-	-	-	-	3-BR	-	-
3.15	A ₂	-	C ₁	-	-	-	-	-	4-F	-	156 - 157
3.16	A ₂	-	C ₁	-	-	-	-	-	4-C ₁	-	142 - 144
3.17	A ₂	-	C ₁	-	-	-	-	-	4-CH ₃	-	115 - 117
3.18	A ₂	-	C ₁	-	-	-	-	-	4-OCH ₃	-	114 - 116



Table 3

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39

No.	A	R ³	R ⁴	R ⁵	R ⁶	R ⁷	R ⁸	R ⁹	R ¹⁰	x	Physical data (°C)
3.19	A ₂	-	C ₁	-	-	-	-	-	4-SCH ₃	-	
3.20	A ₃	-	-	-	-	-	-	2-F	CH ₂		
3.21	A ₃	-	-	-	-	-	-	3-F	CH ₂		
3.22	A ₃	-	-	-	-	-	-	4-F	CH ₂		
3.23	A ₃	-	-	-	-	-	-	3-C ₁	CH ₂		
3.24	A ₃	-	-	-	-	-	-	3-CH ₃	CH ₂		
3.25	A ₃	-	-	-	-	-	-	2-F	S		
3.26	A ₃	-	-	-	-	-	-	3-F	S		
3.27	A ₃	-	-	-	-	-	-	4-P	S		
3.28	A ₃	-	-	-	-	-	-	3-C ₁	S		
3.29	A ₃	-	-	-	-	-	-	3-CH ₃	S		
3.30	A ₃	-	-	-	-	-	-	2-F	SO ₂		
3.31	A ₃	-	-	-	-	-	-	3-F	SO ₂		
3.32	A ₃	-	-	-	-	-	-	4-F	SO ₂		
3.33	A ₃	-	-	-	-	-	-	3-C ₁	SO ₂		
3.34	A ₃	-	-	-	-	-	-	3-CH ₃	SO ₂		
3.35	A ₅	-	-	-	CP ₃	CB ₃	-	2-F	-		
3.36	A ₅	-	-	-	CP ₃	CH ₃	-	3-F	-		
3.37	A ₅	-	-	-	CP ₃	CH ₃	-	4-F	-		
3.38	A ₇	-	-	-	-	-	CH ₃	C ₁	2-F	-	
3.39	A ₇	-	-	-	-	-	CH ₃	C ₁	3-F	-	



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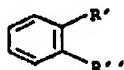
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No.	A	R ³	R ⁴	R ⁵	R ⁶	R ⁷	R ⁸	R ⁹	R ¹⁰	X	Physical data (°C)
3.40	A ₇	-	-	-	-	-	CH ₃	C1	4-F	-	
3.41	A ₇	-	-	-	-	-	CF ₃	C1	2-F	-	
3.42	A ₇	-	-	-	-	-	CF ₃	C1	4-F	-	

THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS:

1. A composition for controlling harmful fungi including, in a solid or liquid carrier,

A) at least one active ingredient I, which inhibits respiration on cytochrome complex III, of the formula



wherein

R' is $-\text{C}(\text{CO}_2\text{CH}_3)=\text{CHOCH}_3$, $-\text{C}(\text{CO}_2\text{CH}_3)=\text{NOCH}_3$, $-\text{C}(\text{CONHCH}_3)=\text{NOCH}_3$, $-\text{C}(\text{CO}_2\text{CH}_3)=\text{CHCH}_3$, $-\text{C}(\text{CO}_2\text{CH}_3)=\text{CHCH}_2\text{CH}_3$, $-\text{C}(\text{COCH}_3)=\text{NOCH}_3$, $-\text{C}(\text{COCH}_2\text{CH}_3)=\text{NOCH}_3$, $-\text{N}(\text{OCH}_3)-\text{CO}_2\text{CH}_3$, $-\text{N}(\text{CH}_3)-\text{CO}_2\text{CH}_3$, or $-\text{N}(\text{CH}_2\text{CH}_3)-\text{CO}_2\text{CH}_3$

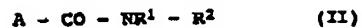
R'' has one of the following meanings:

- a) phenoxymethylen, pyridinyloxymethylene, pyrimidinyloxymethylene or pyrazolylloxymethylene, the aromatic radical being unsubstituted or having 1, 2 or 3 substituents, independently of one another selected from amongst alkyl, halogen, CF_3 , CHF_2 , $-\text{C}(\text{CH}_3)=\text{NOCH}_3$, and phenyl which is unsubstituted or substituted by 1, 2 or 3 halogen atoms and/or alkyl groups;
- b) phenoxy or pyrimidinyloxy, unsubstituted or substituted by 1, 2 or 3 halogen atoms or a phenoxy radical which is unsubstituted or has a halogen or cyano substituent;
- c) $\text{CH}_2\text{ON}=\text{CR}^a\text{R}^b$
where R^a is alkyl; and
 R^b is phenyl which is unsubstituted or has 1, 2 or 3 substituents, independently of one another selected from amongst alkyl, halogen, CF_3 and CHF_2 ;
- d) $\text{CH}_2\text{ON}=\text{CR}^c\text{CR}^d=\text{NOR}^e$
where R^c is alkyl;
 R^d is alkyl or phenyl which is unsubstituted or is



substituted by 1, 2 or 3 halogen atoms; and
 R⁴ is alkyl or alkenyl; and

B) at least one amide compound of the formula II



where

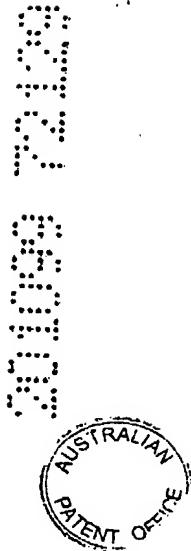
A is one of the following groups:

pyridyl, thiazolyl, pyrazolyl or oxazolyl, it being possible for these groups to have 1, 2 or 3 substituents, independently of one another selected from amongst alkyl, halogen, difluoromethyl and trifluoromethyl,

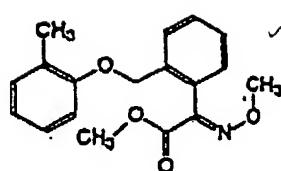
R¹ is a hydrogen atom, alkyl or alkoxy;

R² is a phenyl or cycloalkyl group which has a phenyl substituent and wherein the phenyl substituent may have 1 to 5 halogen atoms and/or 1 to 3 alkyl substituents.

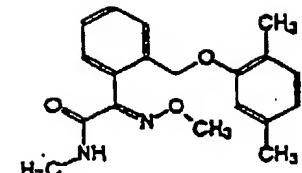
2. A composition as claimed in claim 1 where R⁴ is C(CO₂CH₃)=CHCH₃ or C(CO₂CH₃)=NOCH₃.
3. A composition as claimed in claim 1 which includes, as active ingredient I, a compound of the following formulae:



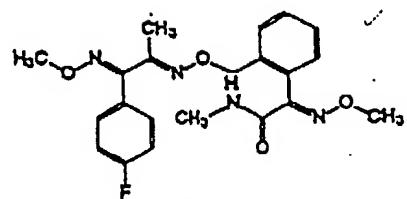
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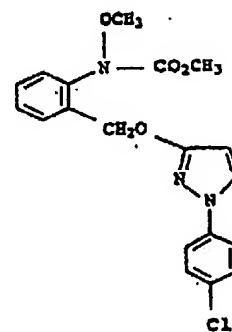
I.3



I.4



I.5



I.8 A-10

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4. A composition as claimed in any of claims 1 to 3 which includes as the amide compound, a compound of the formula II where the radical A is one of the following groups:

pyridin-3-yl which is unsubstituted or substituted in the 2-position by halogen, methyl, difluoromethyl, trifluoromethyl, methoxy, methylthio, methylsulfinyl or methylsulfonyl;

thiazol-5-yl which is unsubstituted or substituted in the 2- and/or 4-position by methyl, chlorine, difluoromethyl or trifluoromethyl;

thiazol-4-yl which is unsubstituted or substituted in the 2- and/or 5-position by methyl, chlorine, difluoromethyl or trifluoromethyl;

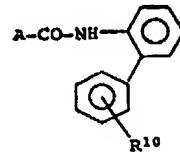
1-methylpyrazol-4-yl which is unsubstituted or substituted in the 3- and/or 5-position by methyl, chlorine, difluoromethyl or trifluoromethyl; or

oxazol-5-yl which is unsubstituted or substituted in the 2- and/or 4-position by methyl or chlorine.

5. A composition as claimed in any one of the preceding claims which includes, as the amide compound, a compound of the formula II where R^2 is a phenyl group which is substituted by the substituent mentioned in claim 1.

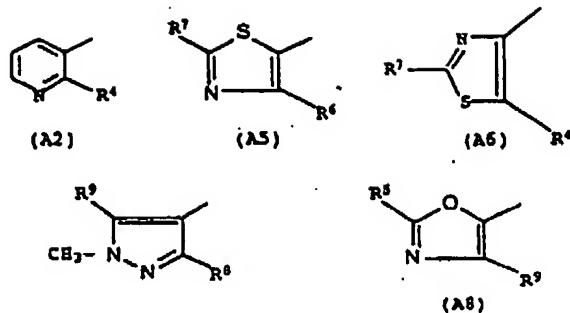
6. A composition as claimed in claim 5 where R^2 is a phenyl group which has the substituent as defined in claim 1 in the 2-position.

7. A composition as claimed in any one of the preceding claims which includes, as the amide compound, a compound of the formula IIa:



where A is

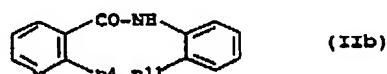




where

R^4 is trifluoromethyl or chlorine,
 R^6 is methyl, difluoromethyl, trifluoromethyl or chlorine,
 R^7 is hydrogen, methyl or chlorine,
 R^8 is methyl, difluoromethyl or trifluoromethyl,
 R^9 is hydrogen, methyl, difluoromethyl, trifluoromethyl or chlorine, and
 R^{10} is C_1-C_4 -alkyl or halogen.

8. A composition as claimed in any of claims 1 to 7 which includes, as the amide compound, a compound of the formula IIb

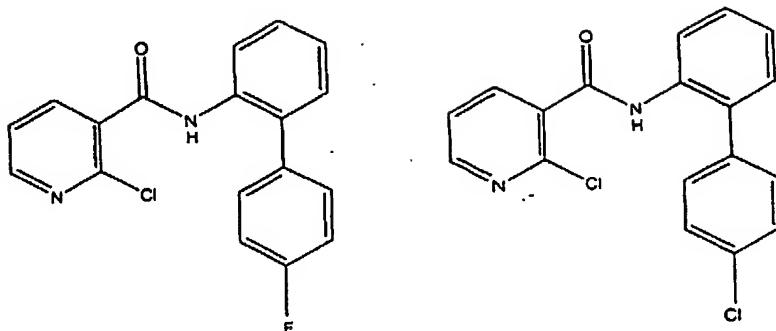


where

R^4 is halogen and
 R^{11} is phenyl which is substituted by halogen.

9. A composition as claimed in any of claims 1 to 8 which includes, as the amide compound, a compound of the formulae below:





10. A method of controlling harmful fungi, which includes treating the fungi, their environment, or the materials, plants, seeds, soils, areas or spaces to be protected against fungal infection, with a composition as claimed in any of claims 1 to 9, it being possible to apply the active ingredients simultaneously, i.e. jointly or separately, or in succession.

11. A two part kit containing:

- A) as a first part thereof, an active ingredient I as defined in any one of claims 1, 2 or 3 in a solid or liquid carrier, and
- B) as the second part thereof, an amide compound of formula II as defined in any one of claims 1, 4, 5, 6, 7, 8 or 9 in a solid or liquid carrier, which first and second parts upon mixing yield a composition as claimed in any one of the preceding claims, the active ingredient I and the amide of formula II being present in proportions which provide synergistic effect in controlling harmful fungi when used in a method according to claim 10.

12. A composition according to claim 1 and as herein described with reference to the examples.



13. A method according to claim 11 and as herein described with reference to the examples.

DATED this 11th day of February, 2000

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